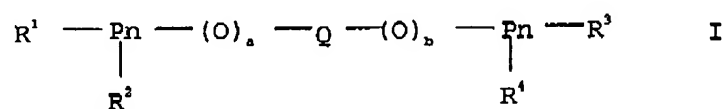


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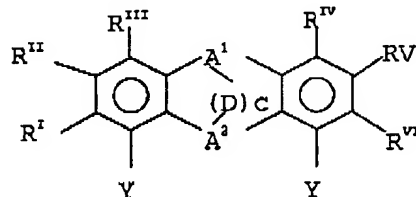
### AMENDMENTS TO THE CLAIMS

1. (Original) A process for preparing dialdehydes and/or ethylenically unsaturated monoaldehydes by reacting at least one compound having at least two ethylenically unsaturated double bonds with carbon monoxide and hydrogen in the presence of a hydroformylation catalyst comprising at least one complex of a metal of transition group VIII with at least one ligand selected from among chelating pnictogen compounds of the formula I,



where

Q is a bridging group of the formula



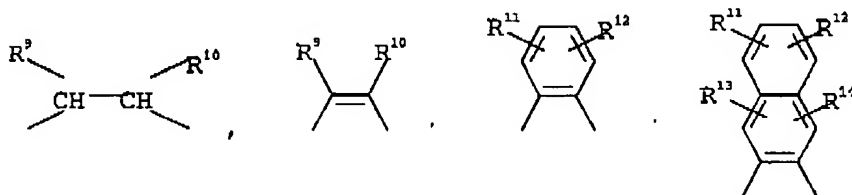
where

A<sup>1</sup> and A<sup>2</sup> are each, independently of one another, O, S, SiR<sup>a</sup>R<sup>b</sup>, NR<sup>c</sup> or CR<sup>d</sup>R<sup>e</sup>, where

R<sup>a</sup>, R<sup>b</sup> and R<sup>c</sup> are each, independently of one another, hydrogen, alkyl, cycloalkyl, heterocycloalkyl, aryl or hetaryl,

$R^d$  and  $R^e$  are each, independently of one another, hydrogen, alkyl, cycloalkyl, heterocycloalkyl, aryl or hetaryl or the group  $R^d$  together with a further group  $R^d$  or the group  $R^e$  together with a further group  $R^e$  form an intramolecular bridging group D,

D is a divalent bridging group selected from among the groups



where

$R^9$  and  $R^{10}$  are each, independently of one another, hydrogen, alkyl, cycloalkyl, aryl, halogen, trifluoromethyl, carboxyl, carboxylate or cyano or are joined to one another to form a  $C_3$ - $C_4$ -alkylene bridge,

$R^{11}$ ,  $R^{12}$ ,  $R^{13}$  and  $R^{14}$  are each, independently of one another, hydrogen, alkyl, cycloalkyl, aryl, halogen, trifluoromethyl,  $COOH$ , carboxylate, cyano, alkoxy,  $SO_3H$ , sulfonate,  $NE^1E^2$ , alkylene- $NE^1E^2E^{3+}X^-$ , acyl or nitro,

c 0 or 1,

Y is a chemical bond,

$R^I$ ,  $R^{II}$ ,  $R^{III}$ ,  $R^{IV}$ ,  $R^V$  and  $R^{VI}$  are each, independently of one another, hydrogen, alkyl, cycloalkyl, heterocycloalkyl, aryl, hetaryl,  $COOR^f$ ,  $COO^-M^+$ ,  $SO_3R^f$ ,  $SO_3^-M^+$ ,

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$NE^1E^2$ ,  $NE^1E^2E^{3+}X^-$ , alkylene- $NE^1E^2E^{3+}X^-$ ,  $OR^f$ ,  $SR^f$ ,  $(CHR^ECH_2O)_xR^f$ ,  
 $(CH_2N(E^1))_xR^f$ ,  $(CH_2CH_2N(E^1))_xR^f$ , halogen, trifluoromethyl, nitro, acyl or cyano,

where

$R^f$ ,  $E^1$ ,  $E^2$  and  $E^3$  are identical or different radicals selected from among hydrogen, alkyl, cycloalkyl and aryl,

$R^E$  is hydrogen, methyl or ethyl,

$M^+$  is a cation,

$X^-$  is an anion, and

x is an integer from 1 to 120,

or

two adjacent radicals selected from among  $R^I$ ,  $R^{II}$ ,  $R^{III}$ ,  $R^{IV}$ ,  $R^V$  and  $R^{VI}$  together with two adjacent carbon atoms of the benzene ring to which they are bound for a fused ring system having 1, 2 or 3 further rings,

a and b are each, independently of one another, 0 or 1,

Pn is a pnictogen atom selected from among the elements phosphorus, arsenic and antimony,

and

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$R^1, R^2, R^3, R^4$  are each, independently of one another, hetaryl, hetaryloxy, alkyl, alkoxy, aryl, aryloxy, cycloalkyl, cycloalkoxy, heterocycloalkyl, heterocycloalkoxy or an  $NE^1E^2$  group, with the proviso that  $R^1$  and  $R^3$  are pyrrole groups bound via the nitrogen atom to the pnicoen atom Pn

or  $R^1$  together with  $R^2$  and/or  $R^3$  together with  $R^4$  form a divalent group E of the formula

Py-I-W

where

Py is a pyrrole group which is bound via the pyrrole nitrogen atom to the pnicoen atom Pn,

I is a chemical bond or O, S,  $SiR^aR^b$ ,  $NR^c$ , substituted or unsubstituted  $C_1$ - $C_{10}$ -alkylene or  $CR^hR^i$ ,

W is cycloalkyl, cycloalkoxy, aryl, aryloxy, hetaryl or hetaryloxy,

and

$R^h$  and  $R^i$  are each, independently of one another, hydrogen, alkyl, cycloalkyl, heterocycloalkyl, aryl or hetaryl,

or  $R^1$  together with  $R^2$  and/or  $R^3$  together with  $R^4$  form a bispyrrole group of the formula

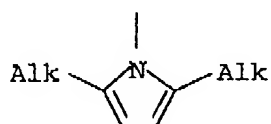
Py-I-Py

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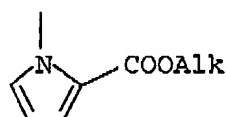
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bound via the nitrogen atoms to the pnictogen atom Pn.

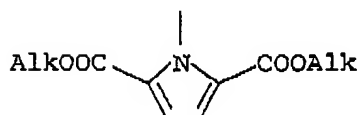
2. (Original) A process as claimed in claim 1, wherein at least one ligand of the formula I, in which the radicals  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are selected independently from among groups of the formulae I.a to I.k



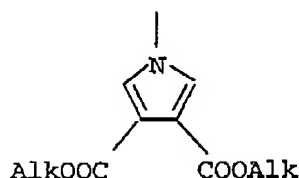
(I.a)



(I.b)



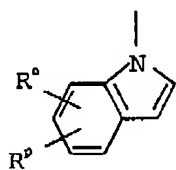
(I.c)



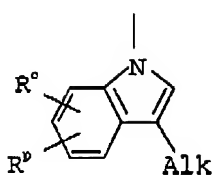
(I.d)

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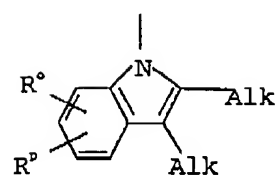
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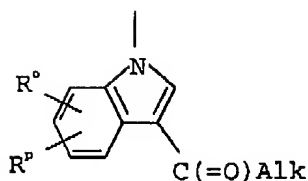
(I.e)



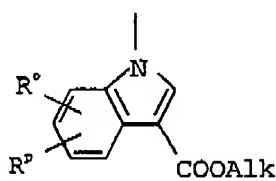
(I.f)



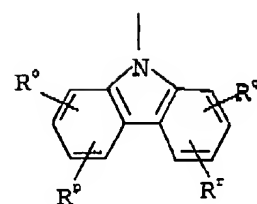
(I.g)



(I.h)



(I.i)



(I.k)

where

Alk is a C<sub>1</sub>-C<sub>4</sub>-alkyl group and

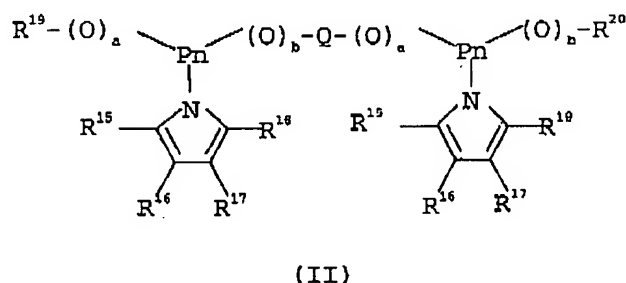
R<sup>o</sup>, R<sup>p</sup>, R<sup>q</sup> and R<sup>r</sup> are each, independently of one another, hydrogen, C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-alkoxy, acyl, halogen, trifluoromethyl, C<sub>1</sub>-C<sub>4</sub>-alkoxycarbonyl or carboxyl, is used.

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3. (Currently amended) A process as claimed in claim 2, wherein at least one ligand of the formula I, in which the radicals  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are each, independently of one another, a 3-alkylindolyl group, ~~preferably a 3-methylindolyl group~~, is used.
4. (Previously presented) A process as claimed in claim 1, wherein the chelating pnictogen compound of the formula I is selected from among chelating pnictogen compounds of the formula II,



where

$R^{15}$ ,  $R^{16}$ ,  $R^{17}$  and  $R^{18}$  are each, independently of one another, hydrogen, alkyl, cycloalkyl, heterocycloalkyl, aryl, hetaryl,  $W'COOR^k$ ,  $W'COO^-M^+$ ,  $W'(SO_3)R^k$ ,  $W'(SO_3)^-M^+$ ,  $W'PO_3(R^k)(R^l)$ ,  $W'(PO_3)^{2-}(M^+)_2$ ,  $W'NE^4E^5$ ,  $W'(NE^4E^5E^6)^+X^-$ ,  $W'OR^k$ ,  $W'SR^k$ ,  $(CHR^lCH_2O)_yR^k$ ,  $(CH_2NE^4)_yR^k$ ,  $(CH_2CH_2NE^4)_yR^k$ , halogen, trifluoromethyl, nitro, acyl or cyano,

where

$W'$  is a single bond, a heteroatom or a divalent bridging group having from 1 to 20 bridge atoms,

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$R^k, E^4, E^5, E^6$  are identical or different radicals selected from among hydrogen, alkyl, cycloalkyl and aryl,

$R^l$  is hydrogen, methyl or ethyl,

$M^+$  is a cation equivalent,

$X^-$  is an anion equivalent and

$y$  is an integer from 1 to 240,

where two adjacent radicals  $R^{15}, R^{16}, R^{17}$  and  $R^{18}$  together with the carbon atoms of the pyrrole ring to which they are bound may also form a fused ring system having 1, 2 or 3 further rings,

with the proviso that at least one of the radicals  $R^{15}, R^{16}, R^{17}$  and  $R^{18}$  is not hydrogen and  $R^{19}$  and  $R^{20}$  are not joined to one another,

$R^{19}$  and  $R^{20}$  are each, independently of one another, cycloalkyl, heterocycloalkyl, aryl or hetaryl, or  $R^{19}$  together with  $R^{15}$  or  $R^{16}$  and/or  $R^{19}$  together with  $R^{17}$  or  $R^{18}$  form a divalent group

-I-W-

where

I is a chemical bond or O, S,  $SiR^aR^b$ ,  $NR^c$  or substituted or unsubstituted  $C_1$ - $C_{10}$ -alkylene, preferably  $CR^hR^i$ , where  $R^a, R^b, R^c, R^h$  and  $R^i$  are each, independently of one another, hydrogen, alkyl, cycloalkyl, heterocycloalkyl, aryl or hetaryl and



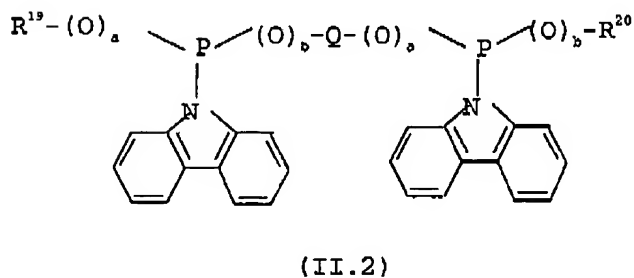
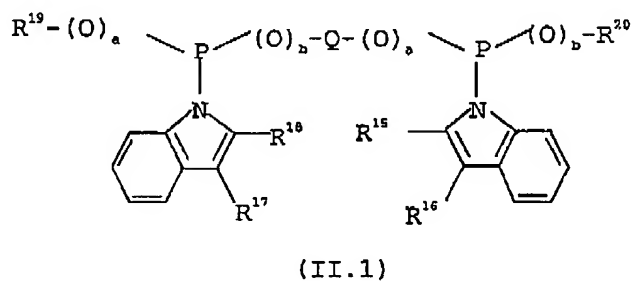
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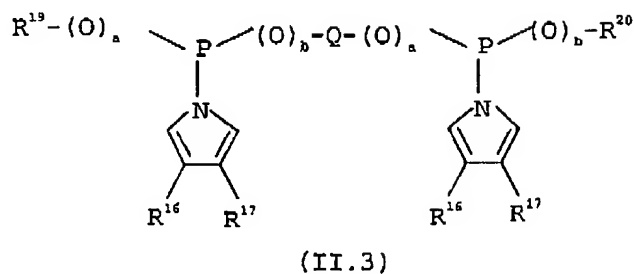
W is cycloalkyl, cycloalkoxy, aryl, aryloxy, hetaryl or hetaryloxy.

5. (Currently amended) A process as claimed in claim 1, wherein the chelating pnictogen compound of the formula I is a chelating pnictogen compound of the formulae II.1 to II.3,



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where

$R^{15}$ ,  $R^{16}$ ,  $R^{17}$ ,  $R^{18}$ ,  $Q$ ,  $a$  and  $b$  are as defined in claim 4;

$Q$ ,  $a$  and  $b$  are as defined in claim 1,

$R^{15}$ ,  $R^{16}$ ,  $R^{17}$  and  $R^{18}$  are each, independently of one another, hydrogen, alkyl, cycloalkyl, heterocycloalkyl, aryl, hetaryl,  $W'COOR^k$ ,  $W'COOM^+$ ,  $W'(SO_3)R^k$ ,  $W'(SO_3)M^+$ ,  $W'PO_3(R^k)(R^k)$ ,  $W'(PO_3)^2-(M^+)_2$ ,  $W'NE^4E^5$ ,  $W'(NE^4E^5E^6)^+X$ ,  $W'OR^k$ ,  $W'SR^k$ ,  $(CHR^1CH_2O)_yR^k$ ,  $(CH_2NE^4)_yR^k$ ,  $(CH_2CH_2NE^4)_yR^k$ , halogen, trifluoromethyl, nitro, acyl or cyano,

wherein

$W'$  is a single bond, a heteroatom or a divalent bridging group having from 1 to 20 bridge atoms,

$R^k$ ,  $E^4$ ,  $E^5$ ,  $E^6$  are identical or different radicals selected from among hydrogen, alkyl, cycloalkyl and aryl,

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R<sup>1</sup> is hydrogen, methyl or ethyl,

M<sup>+</sup> is a cation equivalent,

X<sup>-</sup> is an anion equivalent and

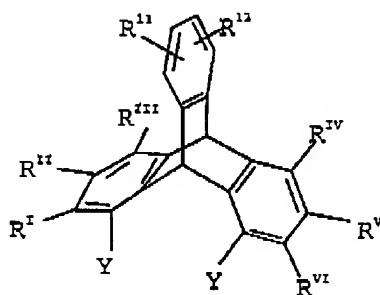
y is an integer from 1 to 240,

where two adjacent radicals R<sup>15</sup>, R<sup>16</sup>, R<sup>17</sup> and R<sup>18</sup> together with the carbon atoms of the pyrrole ring to which they are bound may also form a fused ring system having 1, 2 or 3 further rings,

where at least one of the radicals R<sup>16</sup> and R<sup>17</sup> in the formula II.3 is not hydrogen,

R<sup>19</sup> and R<sup>20</sup> are each, independently of one another, cycloalkyl, heterocycloalkyl, aryl or hetaryl.

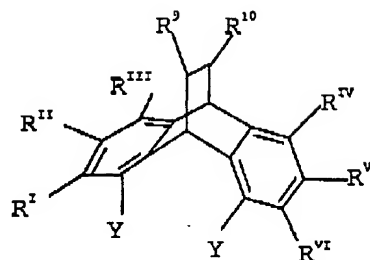
6. (Previously presented) A process as claimed in claim 1, wherein the bridging group Q is a triptycenediyl group of the formula



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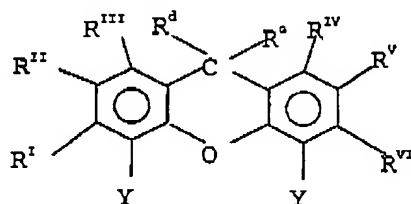
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or the formula



where  $R^I$ ,  $R^{II}$ ,  $R^{III}$ ,  $R^{IV}$ ,  $R^V$  and  $R^{VI}$ ,  $R^9$ ,  $R^{10}$ ,  $R^{11}$  and  $R^{12}$  are as defined in claim 1.

7. (Previously presented) A process as claimed in claim 1, wherein the bridging group Q is a xanthenediyl group of the formula



where  $R^I$ ,  $R^{II}$ ,  $R^{III}$ ,  $R^{IV}$ ,  $R^V$  and  $R^{VI}$  and Y are as defined in claim 1 and  $R^d$  and  $R^e$  are each, independently of one another, hydrogen, alkyl, cycloalkyl, heterocycloalkyl, aryl or hetaryl.

8. (Previously presented) A process as claimed in claim 1, wherein a molar ratio of ligand to metal of transition group VIII of from 1:1 to 1000:1 is set in the reaction mixture.

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9. (Previously presented) A process as claimed in claim 1, wherein the reaction is carried out at from 40 to 80°C.
10. (Previously presented) A process as claimed in claim 1, wherein the compound having at least two ethylenically unsaturated double bonds which is used is a  $\alpha,\omega$ -diolefin.
11. (Previously presented) A process as claimed in claim 1, wherein
  - (i) a compound having a least two ethylenically unsaturated double bonds is subjected to the hydroformylation reaction in a reaction zone,
  - (ii) an output is taken from the reaction zone and is fractionated to give a fraction enriched in unsaturated monoaldehydes and a fraction depleted in unsaturated monoaldehydes, and
  - (iii) the fraction depleted in unsaturated monoaldehydes is recirculated, optionally after work up, to the reaction zone.
12. (New) A process as claimed in claim 2, wherein at least one ligand of the formula I, in which the radicals  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are each, independently of one another, a 3-methylindolyl group.